AMENDMENT UNDER 37 C.F.R. § 1.116

Application No.: 10/662,345

Attorney Docket No.: Q71975

AMENDMENTS TO THE SPECIFICATION

Please add the following new Abstract of the Disclosure:

A method of performing interactive clinical trials for testing a new drug. A pre-clinical

phase is performed in which a computer model for pharmacokinetics and pharmacodynamics of

the drug is created and adjusted based on in vitro studies and in vivo studies in animals. A phase

I clinical research is performed in which a clinical trial on at least a single dose is performed in

parallel with performing computer simulation studies using the computer model. An optimal

protocol is determined for the most responsive patient populations and indications for a phase II

clinical trial. Phase II clinical trial is performed where a number of small scale clinical trials are

performed in parallel based on results of the above. Phase III clinical research is performed for

chosen indications by chosen protocols. Phase IV studies are performed for post-marketing

subpopulation analysis and long term product safety assessment.

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